us-09-978-188a-7.rsp

GenCore version 5.1.6 Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - protein search, using sw model

Run on: April 22, 2004, 13:50:23; Search time 18 Seconds (without alignments) 1423.251 Million cell updates/sec

Title: Perfect score: US-09-978-188A-7 2527

Sequence:

1 MVKFPALTHYWPLIRFLVPL......TDMPPTEEVTDIVEMREENE 492

Scoring table: BLOSUM62 Gapop 10.0 , Gapext 0.5

141681 segs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

																										Result No.
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Q8i8u7 drosophila P37482 bacillus su P79176 gorilla gor	P39981 SACCHAIOMYC P37555 bacillus su P79235 pongo pygma			-	ω	DS0365 allowyces m	mus muscul	sal		P45123 haemophilus			00	σ		arabidops	0	P15578 podospora a	7	P58369 tetraodon n	8	P58367 xenopus lae	P58366 rattus norv	mus n	sapi	Description

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40	95	د. 8	501	_	LYSI CORGL	P35865 corynebacte
41	95	3. 8	511	۳	ALG8 DROME	Q9w3v8 drosophila
42	95	3.8	654	μ	NU5M_RHIST	P50367 rhizopus st
43	94.5	3.7	507	بر	TT12_ARATH	
44	94.5	3.7	518	ч	SP5B BACSU	Q00758 bacillus su
45	94	3.7	421	ы	VG2_BPIKE	P03660 bacteriopha

ALIGNMENTS

ANG PRA RAARARARARARARARARARARARARARARARARA	RESULT
ANNER HUMAN STANDARD; PRT; 492 AA. Q98CJI; Q9NOW2; 28-FEB-2003 (Rel. 41, Created) 28-FEB-2003 (Rel. 41, Last sequence update) 19-CCT-2003 (Rel. 42, Last sequence update) 19-CCT-2003 (Rel. 41, Last sequence update) 19-CCT-2003 (Rel. 42, Last sequence of unidentified humo. 18-CRITER-2045194; PubMed=10894769; 19-CCT-2003 (Rel. 41, Last sequence of unidentified humo. 19-CRITER-2045063; PubMed=10897877; 19-CRITER-2045063; PubMed=10806666; 19-CRITER-2045063; PubMed=108066666; 19-CRITER-20450666; 19-CRITER-2045	27 1

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MIM; 605145; -...
MIM; 123000; -...
GO; GO:0016021;
GO; GO:0019867;
GO; GO:00030504;
GO; GO:0007626;
GO; GO:0007626;
GO; GO:0001501;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      pyrophosphate (PPi), probably functioning as PPi transporter.

SUBCELLULAR LOCATION: Integral membrane protein (Probable).

TISSUE SPECIFICITY: Found in osteoblasts from mandibular bone and from iliac bone; not detected in osteoclastic cells.

DISRASE: Defects in ANKH are the cause of craniometaphyseal dysplasia Jackson type (CMDJ) [MIM:123000]. CMDJ is a rare autosomal dominant skeletal disorder characterized by abnormal bone formation and mineralization in membranous as well as endochondral bones. Progressive tickening of the bones can cause narrowing of cranial formaina and can lead to severe visual and neurological impairment, such as facial palsy and deafness.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Ritter H., Leschik G., Uhlmann K., Mis Goldblatt J., Borochowitz Z.U., Kotzot Braun H.-S., Laing N., Tinschert S.; "Heterozygous mutations in ANKH, the h progressive ankylosis gene, result in Nat. Genet. 28:37-41(2001).
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation—the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entitles requires a license agreement (see http://www.isb-sib.ch/announce/or send an email to license@isb-sib.ch).
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Nuernberg P., Thiele H., Chandler D., Hoehne W., Cunnin
Ritter H., Leschik G., Uhlmann K., Mischung C., Harrop
Goldblatt J., Borochowitz Z.U., Kotzot D., Westermann F
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EMBL; AB046801; BAB13407.1;
EMBL; BC009835; AAH09835.1;
EMBL; BC014526; AAH14526.1;
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GO:0019867; C:outer membrane; TAS.
GO:0030504; F:inorganic diphosphate transporter activity; IDA.
GO:0005315; F:inorganic phosphate transporter activity; IDA.
GO:0007626; F:locomotory behavior; NAS.
                                                                                                                                                                                                                                                                                                                                                                                                                    GO:0030500; P:regulation of bone mineralization; TAS GO:0001501; P:skeletal development; NAS.
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, Baur S.T., Shiang R., Grange D.K., Beighton
Hamersma H., Sellars S., Ramesar R., Lidral A
Raposo do Amaral C.M., Gorlin R.J., Mulliken J
ANKH.
transport; Transmembrane;
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CYTOPLASMIC (POTENTIAL).
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Eukaryota; Metazoa; Chordata; Mammalia; Eutheria; Rodentia;

Craniata; Vertebrata; Euteleostomi;

Muridae; Murinae;

Sciurognathi;

Mus musculus (Mouse)

ANKH MOUSE STANDARD; PRT; 492 AA. 09JHZ2; O35138; O35139; O351

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               VTDIVEMREENE
                                                   LGVHGATLGVGSLLAGFVGESTMVAIAACYVYRKQKKKMENESATEGEDSAMIDMPPTEE
                                                                              FAFAELCVVPLRIFSFFPVPVTVRAHLTGWLMTLKKTFVLAPSSVLRIIVLIASLVVLPY
                                                                                                                                                               WWPLALILATORISRPIVNLFVSRDLGGSSAATEAVAILTATYPVGHMPYGWLTEIRAVY
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492 AA;
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P -> PA (in CMDJ).
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TITLE JOURNAL FEATURES source	REFERENCE AUTHORS TITLE JOURNAL MEDLINE PUBMED REFERENCE AUTHORS	RESULT 3 AB046801 LOCUS DEFINITION ACCESSION VERSION KEYWORDS SOURCE ORGANISM	0	DB Q9 DB Q9	8 8 8 8 8 8	Q
Direct Submission Direct Submission Submitted (03-AUG-2000) Osamu Ohara, Kazusa DNA Research Institute, Submitted (03-AUG-2000) Osamu Ohara, Kazusa DNA Research Institute, Department of Human Gene Research; 1532-3, Yana, Kisarazu, Chiba 292-0812, Japan (E-mail:cdmainfo@kazusa.or.jp, URL:http://www.kazusa.or.jp/huge, Tel:81-438-52-3913, Fax:81-438-52-3914) Location/Qualifiers 13928 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="fj05690" /note="vector:pBluescriptII SK plus"	Mammalia; Eutheria; Primates; Catarrhini; Hominidae, 1 (sites) 1 (sites) Nagase, T., Kikuno, R., Nakayama, M., Hirosawa, M. and Cerediction of the coding sequences of unidentified in XVIII. The complete sequences of 100 new cDNA clones which code for large proteins in vitro DNA Res. 7 (4), 273-281 (2000) 20450683 10997877 2 (bases 1 to 3928) Ohara, O., Nagase, T. and Kikuno, R.	AB046801 Homo sapiens mRNA for KIAA1581 AB046801 AB046801.1 GI:10047236 Homo sapiens (human) Homo sapiens Bukaryota; Metazoa; Chordata;	2881 TTATTTTAGGCTATAATACATTTCCTATTTTCGCATTTTCAATAAATGTCTCTAATACA 2940	2761 CCTTTAAAAAATTATAGACACGGTTCACTAAATTGATTTAGTCAGAATTCCTAGACTGA 2820	2521 TGCCAGGTTGCTGTAGGGTAACTTTTGAAGTAGATATATTACCTGGTTCTGCTATCCTTA 2580 2581 GTCATAACTCTGCGGTACAGGTAATTGAGAATGTACTACGGTACTTCCCTCCACACCAT 2640 2581 GTCATAACTCTGCGGTACAGGTAATTGAGAATGTACTACGGTACTTCCCTCCC	2401 GGCTTGCTTTTCCCTCGCCTTTCCTGAAGGTCGCATTAGAGCGAGGTCACATGGAGCATCC 2460 2461 TAACTTTGCATTTTAGTTTTTACAGTGAACCTGAAGCTTTAAGTCTCATCCAGCATTCTAA 2520
8 8 8 8 8 8	\$ \$ \$ \$ \$ \$	\$ \$ \$ \$ \$ \$	\$ \$ \$ \$ \$	Query Best : Match Qy Db	ORIGIN	gene CDS
0 + 0 + 0 + 0	361 AGGGGCCÀTCGCTGCCGTCTTTCACACACTGATAGCTTATAGTGATTTAGGATACTACAT 420	181 CITGAACCGGGCATTGCTGCTGACGAGGATGCAGATGCAGATGCAGATGCAGCTACGC 240		99.7%; Score 2936.6; DB 9; Length 3928; Similarity 99.9%; Pred. No. 0; 9; Conservative 0; Mismatches 4; Indels 0; Gaps GCTCGGCCCGTCGCCTCCCCGCAGAGTCCCCTCGCGGCAGAATGTGTGTG	/db xref="gi:1004732" /db xref="gi:1004732" /db xref="gi:1004732" /translation="APQPRGGAAAAPEAPPPAASVGPRPARPVAPRLPAESPRGSRC /translation="APQPRGGAAAAPEAPPPAASVGPRPARPVAPRLPAESPRGSRC /WGQPTAGTMVKPPALTHYWPLIRPLVPLGITNIAIDFGEQALMRGIAAVEMDAVEML ASYGLAYSLMKFFTGPMSDFKNVGLVFVNSKRDRTXAVLCMVVAGAIAAVETTLIAYS DLGYYIINKLHHVDESVGSKTRRAPLYLAAFPFWDAMAWHAGILLKHKYSFTVGCAS ISDVIAQVVFVAILLHSHLECREPLLIPILSLYMGALVRCTTLCLGYYKNIHDIIPDR SGPELGGDATIRKHLSFWMPLALTIATORISRFIVNLFVSRDLGGSSAATEAVALITA TYPVGHMPYGMLTEIRAVYPAFDKNNPSNKLVSTSNTVTAAHIKKFTFVCMALSLTLC FVMKWTPNVSEKLIJDIIGVDFAFAELCTVPLRIFSFPVPVTVTAAHIKKFTFVCMALSLTLC FVMKWTPNVSEKLIJDIIGVDFAFAELCTVPLRIFSFPVPVTVTAAHIKACYVYRKQ KKKMENESATEGEDSAMTDMPFTEEVTDIVEMREENE"	,

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	1741 GGGGACCTAGTGAATGGTCTTTÁCTGTTGCTATGTAAAAACAAACGAAACAACTGACTT 1800	1681 GGCCTTGATTTAAAGGTTTCGTGTCAATTCTCTAGCATACTGGGTATGCTCACACTGACG 1740 	1621 CTCTTCCCTCTCCCATCGTATTTTGTTCCCTTTTTTTGTTTG	1561 AGGCACGGACGCCATGGGCACTGCAGGGACGGTCAGTCAG	1501 AGACATGCCTCCGACAGAGGAGGTGACAGACATCGTGGAAATGAGAGAGA	1441 GAAGCAGAAAAAGAAGATGGAGAATTGAGTCGGCCACGGAGGGGGAAGACTCTGCCATGAC 1500	1381 CCTGGCGGGCTTTGTGGGAGAATCCACCATGGTCGCCATCGCTGCGTGCTATGTCTACCG 1440	1321 CAGCCTCGTGGTCCTACCCTACCTGGGGGTGCCACGCTGGGCCTGGGCTGGGCTCCCT 1380	1261 ACTGAAGAAACCTTCGTCCTTGCCCCCAGCTCTGTGCTGCGGATCATCGTCCTCATCGC 1320	1201 CTTCTCCTTCTTCCCAGTTCCAGTCACAGTGAGGGCGCATCTCACCGGGTGGCTGATGAC 1260	1141 GATAGACATCATCGGAGTGGACTTTGCCTTTGCAGAACTCTGTGTTTTTCCTTTTGCGGAT 1200	1081 TCTGTCACTCACGCTCTGTTTCGTGATGTTTTGGACACCCAACGTGTCTGAGAAAAATCTT 1140	1021 GAGCACGAGCACACACTCACGCAGCCCACATCAAGAAGTTCACCTTCGTCTGCATGGC 1080	961 GACGGAAATCCGTGCTGTGTATCCTGCTTTCGACAAGAATAACCCCAGCAACAACTGGT 1020 	901 AGAGGCAGTGGCGATTTTGACAGCCACATACCCTGTGGGTCACATGCCATACGGCTGGTT 960 	841 CAGICGGCCIATIGICAACCICTIIGITICCCGGGACCITIGGTGGCAGITCTGCAGCCAC 900	781 AATAAGAAAGATGCTGAGCTTCTGGTGGCCTTTTGGCTCTAATTCTGGCCACACAGAGAAT 840 	721 CAAGAACATTCACGACATCATCCCTGACAGAAGTGGCCCGGAGCTGGGGGGGAGATGCAAC 780
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	2821 AAGAACCTAAACAAAAAAATATTTAAAGATARAATARTATGCTGTATATGTTAGTTAGT 	761 CC 840 CC	701 AC 780 AC	641 AC 720 AC	581 GI	2521 TGCCAGGTTGCTGTAGGGTAACTTTTGAAGTAGATATATTACCTGGTTCTGCTATCCTTA	61 TA 40 TA	401 GG	2341 TTAAATTGTCACAAAAGCGCATCTCCAGATTCCAGACCCTGCCGCATGACTTTTCCTGAA 	CI C	21 AA 00 AA	161 GF 240 GF	AA	2041 GAGCCCGGTGGCCTCTTAAATTTCCCTTCTGCCACGGAGTTCGAAACCATCTACTCCAC 	1981 CAGGTTAAAACTCGGCTTCCTTTGATTTTGCTTCCCAGTCACATGGCCGTACAAAAAGAATG 		1861 TCCTCCCCTGGACAATCTCCTCTTGGAACCAAAGGACTGCAGCTGTGCCATCGCGCCTCG	80 (2)

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              TITLE
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                                                              source
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Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,

Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,

Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,

Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,

Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,

Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,

Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,

Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J.,

McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,

Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,

Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,

Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S.,

Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y.,

Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D.,

Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M.,

Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,

Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.

Generation and initial analysis of more than 15,000 full-length

human and mouse cDNA sequences
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                                                                     through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Series: IRAL Plate: 14 Row: g Column: 4
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 21536393, 100211
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Homo sapiens ankylosis, progressive homolog (mouse), clone MGC:11142 IMAGE:3837372), complete cds.
                                                                                                                                                                                                                                                                                                                          cDNA Library Preparation: Rubin Laboratory cDNA Library Arrayed by: The I.M.A.G.E. Consortium DNA Sequencing by: Institute for Systems Biology
                                                                                                                                                                                                                                                                                                                                                                                              Contact: MGC help desk
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: DCTD/DTP
                                                                                                                                                                                                                                                                                                                                                                                                                                                              NIH-MGC Project URL: http://mgc.nci.nih.gov
On Aug 19, 2003 this sequence version replaced gi:15778895
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Direct Submission
Submitted (24-SEP-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Homo sapiens
                                                                                                                                                                                                                                  http://www.systemsbiology.org
contact: amadan@systemsbiology.org
Anup Madan, Jessica Fahey, Erin Helton,
Madan, Stephanie Rodrigues, Amy Sanchez
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Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
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Best Local Similarity
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Query Match
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